

COMPOSITE POLYMER DIELECTRIC FILM

Background

Integrated circuits contain many different layers of materials, including dielectric layers that insulate adjacent conducting layers from one another. With each decrease in the size of integrated circuits, the individual conducting layers within the integrated circuits grow closer to adjacent conducting layers. This necessitates the use of dielectric layers made of materials with low dielectric constants to prevent problems with capacitance, cross talk, etc. between adjacent conducting layers.

Low dielectric constant polymers have shown promise for use as dielectric materials in integrated circuits. Examples of low dielectric constant polymers include, but are not limited to, fluoropolymers such as TEFLON $((-\text{CF}_2-\text{CF}_2-)_n; k_d = 1.9)$ and PPX-F $((-\text{CF}_2-\text{C}_6\text{H}_4-\text{CF}_2-)_n; k_d = 2.23)$. Many of these materials have been found to be dimensionally and chemically stable under temperatures and processing conditions used in later fabrication steps, have low moisture absorption characteristics, and also have other favorable physical properties.

However, many low dielectric constant polymers have been found to adhere poorly to silicon-containing layers that are commonly used in integrated circuits, including but not limited to silicon oxide, silicon nitride, silicon carbide, and $\text{SiO}_x\text{C}_y\text{H}_z$. Furthermore, the syntheses used to create these films may leave many unreacted free radical chain ends, which may be susceptible to contamination by water, oxygen, and other materials that may reduce the dimensional and chemical stability of the film under

increased temperatures. These problems may result in unreliable device fabrication and low device yields.

Summary

An integrated circuit including a composite polymer dielectric layer formed on a substrate is provided, wherein the composite polymer dielectric layer includes a first silane-containing layer formed on the substrate, wherein the first silane-containing layer is formed at least partially from an organosilane material, a polymer dielectric layer formed on the first silane-containing layer, and a second silane-containing layer formed on the polymer dielectric layer. In some embodiments, the first silane-containing layer and second silane-containing layer may be formed from organosilane materials having at least one unsaturated bond capable of free radical polymerization. Systems and methods for making the disclosed integrated circuits are also provided.

Brief Description of the Drawings

Fig. 1 is a greatly magnified sectional view of an embodiment of a composite polymer dielectric film, formed on an underlying substrate.

Figs. 2a – 2f show chemical reactions that may occur between the underlying substrate and adhesion promoter layer, within the adhesion promoter layer, and between the adhesion promoter layer and polymer dielectric layer of the embodiment of Fig. 1.

Fig. 3 is a flow diagram showing an embodiment of a method of forming a composite polymer dielectric layer on a substrate.

Fig. 4 is a flow diagram showing another embodiment of a method of forming a polymer dielectric layer on a substrate.

Fig. 5 is a block diagram of an exemplary embodiment of a system for forming a composite dielectric layer on a substrate.

Fig. 6 is a block diagram of another exemplary embodiment of a system for forming a composite dielectric layer on a substrate.

Fig. 7 is a plan view of an exemplary embodiment of a system for forming a composite dielectric layer on a substrate.

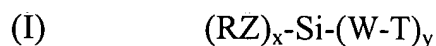
Detailed Description of the Depicted Embodiments

Fig. 1 shows, generally at 10, a first exemplary embodiment of a composite polymer dielectric film formed on a substrate 12. Composite polymer dielectric film 10 includes a first silane-containing adhesion promoter layer 14 disposed on substrate 12, a low dielectric constant polymer layer 16 disposed on first adhesion promoter layer 14, and a second silane-containing layer 18 disposed on low dielectric constant polymer layer 16. Composite polymer dielectric film 10 is typically used as an interlayer in an integrated circuit. Thus, other layers (not shown for purposes of clarity) are typically formed over second silane-containing layer 18. While layer 18 is described in some of the depicted embodiments as being a second adhesion promoter layer, it will be appreciated that layer 18 also may be a hard mask layer or an etch stop layer, such as those used in single and dual damascene processes. Each of these variations is described in more detail below.

As described above, low dielectric constant polymer layer 16 may be formed from materials such as TEFLON, PPX-F and other similar materials. While possessing a low dielectric constant, these materials may not adhere to underlying layers with enough force

to withstand the rigors of downstream processing steps. For example, when deposited directly on a layer of silicon oxide or silicon carbide (which are commonly used in integrated circuits), PPX-F adheres to the layer with a yield strength of approximately 2-3 MPa, indicating that the adhesion of the PPX-F film to the silicon oxide or carbide layer is primarily due to Van der Waals interactions. PPX-F may adhere to other silicon-containing layers with similar yield strengths. In contrast, a film should adhere to an underlying or overlying layer with a yield strength of approximately 4-5 MPa for the structure to safely withstand subsequent back-end processing steps. The use of a properly selected adhesion promoter layer 14 may help to increase the yield strength of the low dielectric constant polymer layer 16 on an underlying or overlying silicon-containing layer sufficiently to equal or surpass a desired yield strength threshold without raising the dielectric constant of the overall composite polymer dielectric film 10 an undesirable amount.

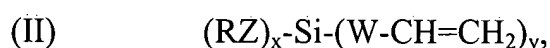
First adhesion promoter layer 14 is formed from a material capable of forming chemical bonds to both substrate 12 and low dielectric constant polymer layer 16. One example of a suitable class of materials to use to form first adhesion promoter layer 14 are silanes having a general structure of



wherein W is selected from the group consisting of -O-, -CH₂-, -(CH₂)_aC=OO-, and -(CH₂)_a-OO=C-; wherein T is selected from the group consisting of -CR=CR'R'', an alkyl halide, and -RC=O; wherein Z is selected from the group consisting of O and NR; wherein R, R' and R'' are an H, alkyl or aromatic group; wherein a is 0 or an integer;

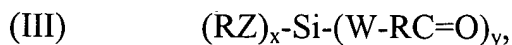
wherein $x = 1, 2$ or 3 ; wherein $y = 1, 2$ or 3 ; and wherein $x + y = 4$. Alternatively, ZR can be H. Such materials are capable of forming strong Si-Z-Si chemical bonds ($Z = O$ or NR, wherein R is an alkyl or aromatic group) to an underlying silicon-containing substrate.

5 A more specific example of compounds of the general formula (I) include vinyl compounds having the general formula



wherein W may be selected from -O- (such as a vinyl ether) -CH₂-, -(CH₂)_aCOO-(such as a vinylacetyl-) and -(CH₂)_aOOC- (such as a vinylacrylo-), wherein Z is selected from the
10 group consisting of O and NR, wherein R is an alkyl or aromatic group, wherein a is zero or an integer, wherein $x = 1, 2$ or 3 , wherein $y = 1, 2$ or 3 , and wherein $x + y = 4$. The vinyl group of these compounds is able to undergo a free radical polymerization with vinyl groups on adjacent vinyl silane molecules, thus helping to form a robust, polymerized adhesion layer that is chemically bonded to the adjacent silicon-containing
15 layer. Where $y = 1$ in structure (II) above, a linear vinyl polymer with side silane groups bonded to the underlying silicon layer is produced. Where $y = 2$ or 3 , a cross-linked vinyl polymer is produced.

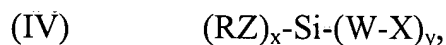
Another specific example of a suitable class of materials of general formula (I) for forming adhesion promoter layer 14 are keto-containing silanes ("Keto silanes") having a
20 general structure of



wherein R is an alkyl group, wherein W is CH₂ or (CH₂)_a, and wherein a is an integer from 1 to 5.

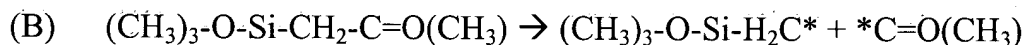
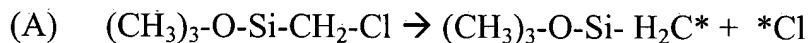
Yet another example of a suitable class of materials of general formula (I) to use to form adhesion promoter layer 14 are halide-containing silanes ("Halide-silanes")

5 having a general structure of

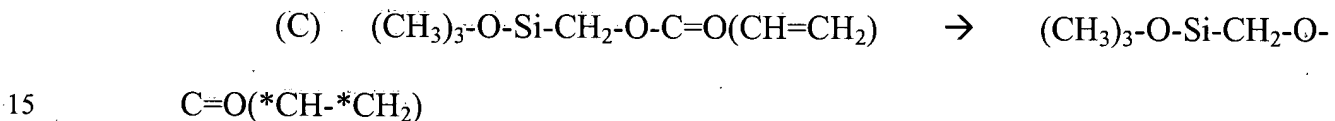


wherein X is a halide, typically Cl, Br or I.

All the above-mentioned vinyl, keto and halide silanes can undergo free radical polymerization when initiated by a suitable energy source, such as UV radiation. For
10 example, under UV irradiation, the following reactions can happen:



and



Furthermore, some free radicals in first adhesion promoter layer 14 may not react with other silane molecules, and instead may undergo a free radical reaction with low dielectric constant polymer layer 16. Synthesis of low dielectric constant polymer layer 16 (described in more detail below) may be carried out by transport polymerization of a
20 gas phase diradical monomer species that is able to polymerize upon condensing on a substrate surface from the gas phase. During transport polymerization, some diradical monomers may react with unreacted vinyl silane-, keto silane- or halide silane-derived

free radicals in first adhesion promotion layer 14, thus chemically bonding first adhesion promoter layer 14 with low dielectric constant polymer layer 16. Thus, by chemically bonding to both substrate 12 and low dielectric constant polymer layer 16, adhesion promoter layer 14 greatly improves the adhesion of the low dielectric constant polymer layer to the substrate.

Figs. 2a-2f illustrate an exemplary series of chemical reactions between the acrylosilane $(C_2H_5O)_3Si(CH_2)_xOOC-CH=CH_2$, an underlying silicon-containing substrate, and an overlying low dielectric constant layer of PPX-F formed from the deposition of the $*CF_2C_6H_4CF_2*$ diradical monomer. First referring to Fig. 2a, the silane functional group of the silane molecule may react with surface oxygen typically found on the surface of a silicon-containing substrate via an elimination of an OCH_2CH_3 leaving group. Here, the surface oxygen in the silicon-containing substrate is shown as being bonded to a hydrogen atom. This bond may be a hydrogen bond, and the hydrogen atom may be part of a water molecule, or the OH may be a hydroxyl group bonded to substrate 12. In either case, the reaction of the silane group with the surface oxygen results in the formation of a chemical bond between the surface oxygen and silicon atom of the silane group, and the production of ethanol from the ethoxide leaving group. In the depicted embodiment, the formation of a single Si-O-Si bond is shown. However, it will be appreciated that more than one Si-O-Si (or, more generally, a Si-Z-Si) bond may form, potentially up to the number of (RZ) leaving groups on the silicon atom in the silane.

Once the silane has been deposited on substrate 12, the silane layer may be exposed to a suitable energy source, including but not limited to heat, UV light or a

plasma, to generate free radicals on the silane molecules. This permits the silane molecules to polymerize to form adhesion promoter layer 14, as depicted in Fig. 2b. Furthermore, as depicted in Fig. 2c, some free radicals in adhesion promoter layer 14 may react with free radicals in low dielectric constant polymer layer 16 during deposition of the low dielectric constant polymer layer, thus chemically bonding first adhesion promoter layer 14 to low dielectric constant polymer layer 16, as shown in Fig. 2d.

After completing the deposition of low dielectric constant polymer layer 16, second adhesion promoter layer 18 may be deposited onto the low dielectric constant polymer layer and exposed to a suitable energy source to generate free radicals from silane molecules in the second adhesion promoter layer (not shown in Figs. 2a-2f) in the same manner as shown for first adhesion promoter layer 14. This permits the silane molecules in the second adhesion promoter layer to polymerize, thus forming a strong polymer layer. Furthermore, this permits some radicals in second adhesion promoter layer 18 to react with unreacted free radicals in low dielectric constant polymer layer 16 to bond the second adhesion promoter layer to the low dielectric constant polymer layer with strong covalent bonds. The result is that the layers of composite dielectric film are chemically bonded together, and that the surfaces of the composite dielectric film include many pending silane groups capable of forming strong chemical bonds to adjacent silicon-containing (or other) layers.

In some embodiments, composite polymer dielectric film 10 may be annealed after formation. Examples of suitable annealing processes are described in more detail below. Annealing composite polymer dielectric film 10 may help to improve the crystallinity and

phase stability of the low dielectric constant polymer layer, as described in U.S. Published Patent Application Serial No. US2003/0036617 to Chung Lee, filed August 9, 2001, the disclosure of which is hereby incorporated by reference. Furthermore, composite polymer dielectric film 10 may be annealed in the presence of hydrogen gas to cap any unreacted free radicals within the adhesion promoter and low dielectric constant polymer layers with hydrogen atoms, thus helping to prevent unwanted reactions with water, oxygen or other free radical scavengers after the composite film on the substrate is removed from the deposition chamber and exposed to ambient.

Annealing composite polymer dielectric film 10 may also cause various chemical changes to occur to the first and second adhesion promoter layers. Some examples of possible chemical changes are shown in Fig. 2e. For example, the hydrocarbon chain disposed between the silane functional group and the acrylo functional group may decompose into gaseous hydrocarbons at a temperature of around 300 degrees Celsius. The CO₂ group within the acrylo functional group and the terminal *CH₂ radical may decompose around 250 degrees Celsius. The ethoxide functional groups may leave as ethanol at 300 to 350 degrees Celsius. The loss of these various functional groups during an annealing process offers the advantage that oxygen is removed from the first and second adhesion promoter layers. The removal of the oxygen at this stage in a device fabrication process may help to prevent problems caused by the oxygen in later device fabrication stages, including but not limited to the formation of bubbles in other layers by oxygen released by later heating stages. However, the loss of these functional groups may cause some of the direct covalent linkages between low dielectric constant polymer

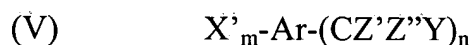
film 16 and substrate 12 (via first adhesion promoter layer 14) to be broken. For example, before annealing composite polymer dielectric film 10, the composite dielectric film has a yield strength of approximately 50 MPa, which is the tensile strength of the composite polymer dielectric film and is more than an order of magnitude greater than the yield strength in the absence of first adhesion promoter layer 14 (2-3 MPa). After annealing, the yield strength of the composite polymer dielectric film may decrease to around 10 MPa. However, this is still well above the yield strength required for downstream processes, which is around 4-5 MPa.

The significant yield strength found after annealing may be attributed to the formation of some covalent bonding formed during the annealing process by the decomposition reactions shown in Fig. 2e. Fig. 2f shows an example of a possible structure that links low dielectric constant polymer layer 16 to substrate 12 via covalent bonding in first adhesion promoter layer 14. It will be appreciated that the structure depicted in Fig. 2f is merely exemplary, and that other structures may also form during an annealing process.

First adhesion promoter layer 14 and second adhesion promoter layer 18 may be formed from any suitable vinyl, keto and/or halide silane materials, including but not limited to those of the general structures (I) - (IV) set forth above. Silanes of the general structure $(RZ)_x-Si-((CH_2)_3-O-C=O(CH=CH_2)_y$, have a high sensitivity to UV irradiation, and thus allow a shorter exposure time to be used in the production of composite film 10. Other silanes, such as acetylsilanes and vinyl ether silanes, may require a higher energy level of UV irradiation, and therefore may require longer exposure times.

First adhesion promoter layer 14 may be formed from the same silane as second adhesion promoter layer 18, or may be formed from a different silane. Furthermore, while composite polymer dielectric film 10 is depicted as having both a first adhesion promoter layer 14 and a second adhesion promoter layer 18, it will be appreciated that the composite dielectric layer may have only one or the other adhesion promoter layer, depending upon whether the adjacent layers in the integrated circuit are able to bond to the low dielectric constant polymer layer 12 with sufficient strength in the absence of an adhesion promoter layer.

Low dielectric constant polymer layer 16 may also be made from any suitable material, including but not limited to poly(paraxylylene) and fluorinated poly(paraxylylene) and derivatives thereof, and other polyaromatics and fluorinated polyaromatics prepared from a monomer of the general structure.



In this formula, X' and Y are leaving groups that can be removed to form a free radical for each removed leaving group, Ar is an aromatic group or a fluorine-substituted aromatic group bonded to m X' groups and n CZ'Z''Y groups, and Z' and Z'' are H, F or C₆H_{5-x}F_x (x = 0, or an integer between 1 and 5). For example, where m = 0 and n = 2, removal of the leaving group y from each CZ'Z''Y functional group yields the diradical Ar(CZ'Z''*)₂. Compounds in which Z' and Z'' are F may have lower dielectric constants and improved thermal stability. Examples of suitable leaving groups for X' and Y include, but are not limited to, ketene and carboxyl groups, bromine, iodine, -NR₂, -N⁺R₃, -SR, -SO₂R, -OR, =N⁺=N-, -C(O)N₂, and -OCF-CF₃ (wherein R is an alkyl or aromatic

group). The numbers m and n in formula (V) may independently be either zero or an integer, and (n + m) is equal to or greater than two, but no greater than the total number of sp² hybridized carbons in the aromatic group that are available for substitution.

Ar in formula (V) may be any suitable aromatic group. Examples of suitable aromatic groups for Ar include, but are not limited to, the phenyl moiety C₆H_{4-n}F_n (n = 0 to 4); the naphthenyl moiety C₁₀H_{6-n}F_n (n = 0 to 6); the di-phenyl moiety C₁₂H_{8-n}F_n (n = 0 to 8); the anthracenyl moiety C₁₂H_{8-n}F_n (n = 0 to 8); the phenanthrenyl moiety C₁₄H_{8-n}F_n (n = 0 to 8); the pyrenyl moiety C₁₆H_{8-n}F_n (n = 0 to 8); and more complex combinations of the above moieties such as C₁₆H_{10-n}F_n (n = 0 to 8). Isomers of various fluorine substitutions on the aromatic moieties are also included. More typically, Ar is C₆F₄, C₁₀F₆, or C₆F₄-C₆F₄.

Low dielectric constant polymer film 16 may also be made from a precursor having the general formula



wherein X' and X'' are leaving groups, and Ar is an aromatic or fluorine-substituted aromatic. The numbers m and n each may be zero or an integer, and m + n is at least two, but no greater than the total number of sp² hybridized carbon atoms on Ar that are available for substitution. For example, polyphenylene (-(C₆H₄)-) and fluorine-substituted versions thereof may be formed from a precursor having general formula (VI). Removal of the leaving groups X' and/or X'' may create the diradical benzyne (*C₆H₄*), which can then polymerize to form polyphenylene. Other aromatic groups besides the phenyl moiety that may be used as Ar in formula (VI) include, but are not

limited to, the naphthenyl moiety $C_{10}H_{6-n}F_n$ ($n = 0$ to 6); the diphenyl moiety $C_{12}H_{8-n}F_n$ ($n = 0$ to 8); the anthracenyl moiety $C_{14}H_{10-n}F_n$ ($n = 0$ to 8); the phenanthrenyl moiety $C_{14}H_{8-n}F_n$ ($n = 0$ to 8); the pyrenyl moiety $C_{16}H_{8-n}F_n$ ($n = 0$ to 8); and more complex combinations of the above moieties such as $C_{16}H_{10-n}F_n$ ($n = 0$ to 10).

5 First adhesion promoter layer 14, low dielectric constant polymer layer 16, and second adhesion promoter layer 18 may each have any suitable thickness. Depending upon the materials selected for each layer, the material or materials from which first and second adhesion promoter layers 14 and 18 are formed may have a higher dielectric constant than the material from which low dielectric constant polymer layer 16 is formed.

10 In this instance, it may be desirable to utilize relatively thin adhesion promoter layers 14 and 18 compared to the low dielectric constant layer 16. For example, in some embodiments, the thickness of first adhesion promoter layer 14 may range from one molecular layer to approximately 50 angstroms, the thickness of low dielectric constant polymer layer may range from one molecular layer up to 3000 angstroms, and the

15 thickness of second adhesion promoter layer 18 may range from one molecular layer to two hundred angstroms. In the specific case of a PPX-F low dielectric constant layer with a thickness of 1500-3000 angstroms, a first adhesion promoter layer with a thickness of about 10 to 30 angstroms, and a second adhesion promoter layer with a thickness of about 200 to 500 angstroms, the overall dielectric constant of the composite dielectric

20 film is approximately 2.4. It will be appreciated that these thickness ranges are merely exemplary, and that first adhesion promoter layer 14, polymer dielectric layer 16 and second adhesion promoter layer 18 may have any other suitable thicknesses.

Instead of serving as an adhesion promoter layer, second adhesion promoter layer 18 may instead (or additionally) be configured to serve as an etch stop for a later etching process in the fabrication of an integrated circuit, or as a hard mask layer in a single or dual damascene process. Where second adhesion promoter layer 18 is used as an etch stop layer, it may be desirable to use a somewhat thicker second adhesion promoter layer 18 than where the second adhesion promoter layer is not used as an etch stop layer.

Where layer 18 is used as a hard mask layer in a damascene process, layer 18 should etch via a similar chemical process, and at similar rates, as low dielectric constant polymer layer 16. In this case, it may be advantageous for layer 18 to have a lower silicon content. In terms of structure (I) where the silane is an ethoxy silane, a compound of the composition $(\text{ETO})\text{-Si-(WT)}_3$ may be used as the hard mask to make the etching characteristics of layer 18 more similar to those of low dielectric constant polymer layer 16 so that the two layers can be etched by similar chemical processes. Examples of suitable etching processes for these two layers include, but are not limited to, O_2 , H_2/N_2 , and NH_3 etching processes.

On the other hand, where layer 18 is used as an etch stop layer, the silicon content of layer 18 may be increased so that layer 18 is etched at a substantially slower rate than low dielectric constant polymer layer 16. In terms of equation (I) where the silane is an ethoxy silane, layer 18 may have a composition of $(\text{ETO})_3\text{-Si-(WT)}$ to give layer 18 a substantially slower etching rate than layer 16 for O_2 , H_2/N_2 , and NH_3 etching processes. On the other hand, a high silicon-content layer 18 may be etched relatively quickly with etching chemistries used for silicon, such as an SF_4 etching process.

Composite polymer film 10 may be formed in any suitable manner. Fig. 3 shows, generally at 100, one exemplary method of forming composite polymer dielectric film 10. Method 100 includes, at 102, first exposing a substrate on which the composite film is to be formed to UV light. Exposing the substrate to UV light removes water from the substrate surface, and helps first adhesion promoter layer 14 to adhere more strongly to the substrate surface. Prior methods of removing water from the surface of the substrate include drying the wafer surface with a desiccant, and with a combination of desiccation and heat. It has been found that exposing the substrate to UV light for just a few second helps to achieve the same improvements in adhesion as desiccating the substrate for approximately eight hours, or desiccating and heating the substrate for approximately six hours. Furthermore, it has been found that PPX-F deposited directly onto the surface of the substrate (without use of first adhesion promoter layer 14) after exposing the substrate to UV light has a yield strength of approximately 3-4 MPa. This is a sufficiently high yield strength to withstand integrated circuit later processing steps. However, the use of first adhesion promoter layer 14 provides a much higher yield strength, and therefore provides a much larger margin of safety in a device fabrication process.

After exposing the substrate to UV light at 102, method 100 next includes depositing a first layer of silane onto the substrate at 104. The silane may be a material configured to chemically bond both to the substrate and to a later-deposited low dielectric constant polymer film, including but not limited to the vinyl, keto and halide silanes listed above. The silane may be deposited in any suitable manner. For example, the silane may be deposited by flowing an inert carrier gas, such as nitrogen, containing a

concentration of the silane across the surface of the substrate. Alternatively, the first layer of silane may be applied by spin-on coating of a solution of the silane in a solvent such as isopropyl alcohol. The substrate may be held at any suitable temperature while the silane is deposited. In some embodiments, the substrate is held between approximately zero and -20 degrees Celsius during deposition of the silane in order to increase the adsorption rate and increase system throughput for the deposition of composite film 10. Furthermore, the first layer of silane deposited at 104 may be deposited in a layer of any desired thickness. For example, the silane may be deposited to a thickness of between one molecular layer and approximately 30 angstroms, or may be deposited in a layer thicker than 30 angstroms where suitable.

After depositing the first layer of silane at 104, the first layer of silane is exposed to a free radical-generating energy source, such as UV light, at 106. Exposure to the free radical-generating energy source at 106 generates free radicals from the silane, and may initiate polymerization of the first adhesion promoter layer, for example, if the silane is an acrylo silane. The UV lamp may output UV light of any suitable wavelength, including but not limited to wavelengths between 170 nm and 350 nm. More specifically, wavelengths between 220 and 300 nm may optionally be employed to help to avoid the formation of ozone.

If desired, the amount of UV energy to which the first silane layer is exposed may be limited to only partially polymerize the layer. This may result in the formation of a larger number of unreacted CH^* and CH_2^* radicals for reacting with a subsequently-formed low dielectric constant polymer layer. While the free radical-generating energy

source is depicted in Fig. 3 as being a UV lamp, it will be appreciated that any other suitable type of energy source may be used. Examples include, but are not limited to, thermal sources, especially if the silane is a silane. In either case, exposure to the free radical-generating energy source may be performed in the absence of free radical scavengers, including but not limited to oxygen, water, and amino-containing compounds. This may help to avoid growing polymer chains from being terminated by the free radical scavengers. The resulting silane polymers include many pending $\text{Si}(\text{ZR})_x$ groups that may bond to the underlying silicon-containing substrate, and also free radical chain ends $-\text{H}_2\text{C}^*$ capable of reacting with low dielectric constant polymer layer 16.

Next, low dielectric constant polymer film 16 is formed by depositing, at 108, a gas-phase diradical monomer onto first adhesion promoter layer 14. During this process, the substrate may be cooled sufficiently to cause the gas phase diradical monomer molecules to condense on first adhesion promoter layer 14. However, the substrate should not be cooled to such an extent that the gas phase diradical monomer molecules to immediately stick to adhesion promoter layer 14 in any orientation. Instead, the substrate should be cooled to a temperature that allows the gas phase diradical monomer molecules to have sufficient energy to “bounce” on the surface of the first adhesion promoter layer so that the monomer diradical molecules can find an energetically favorable orientation. This may allow the low dielectric constant polymer layer to grow with a relatively high degree of crystallinity. Suitable temperature ranges for achieving this effect may differ depending upon the monomer being deposited. However, in general, temperatures below the melting temperature of the diradical intermediates or the growing low dielectric

constant polymers are suitable for achieving this effect. For the diradical monomer $^*\text{CF}_2\text{C}_6\text{H}_4\text{CF}_2^*$, suitable temperatures may include those between approximately -30 and -50 degrees Celsius.

It has been found that, with the temperature held at a suitable level, PPX-F can be deposited from the gas phase diradical monomer $^*\text{CF}_2\text{C}_6\text{H}_4\text{CF}_2^*$ with an initial crystallinity of 20-50% in a β_2 phase of the material. Furthermore, it has been found that the polymer chains having the 20-50% initial crystallinity have a high degree of alignment along a single crystallographic direction, and that the crystallinity can thus be increased to 70% simply by annealing the film at a temperature higher than the glass transition temperature (approximately 170 degrees Celsius for PPX-F), but lower than the melting temperature of the polymer, for intervals as short as three minutes or less. The low dielectric constant polymer film formed at 108 may be annealed before the second layer of silane material is deposited, or may be annealed after all layers of composite polymer dielectric film 10 have been deposited, as described below.

Next, the second layer of silane is deposited onto the polymer dielectric layer at 110. This layer is then exposed to UV light at 112 to initiate polymerization of the silane molecules to form the second adhesion promoter layer 18 (or an etch stop or hard mask layer), and to react the second adhesion promoter layer, the etch stop or the head mask layer with low dielectric constant polymer layer 16. The second silane layer may be deposited in any suitable thickness. As described above, where second silane layer 18 is to be used as an etch-stop or hard mask layer, the second silane layer may be applied in a thicker layer ranging from 200 to 500 angstroms. Likewise, the second silane layer may

be applied in a thinner layer ranging from 5 to 20 angstroms when layer 18 is not to be used as an etch-stop or hard mask layer to reduce the impact of the second adhesion promoter layer on the overall dielectric constant of the structure.

The second silane layer may be deposited in any suitable manner, including but not limited to vapor phase deposition techniques, either with or without a carrier gas, and spin-on deposition techniques utilizing a solution of silane in a suitable solvent.

After depositing the second layer of silane onto the low dielectric at 110 and exposing the second layer of silane to UV light (or other free radical-generating energy source) at 112, the composite polymer dielectric film may be annealed in the presence of a reducing gas, such as hydrogen, at 114. After formation of the composite polymer dielectric film, many unreacted free radicals $\cdot\text{CH}$, $\cdot\text{CH}_2$, and $\cdot\text{CF}_2$ may remain in the layers making up the composite film. Annealing in the presence of hydrogen may eliminate these free radicals before composite polymer dielectric film 10 is removed from the deposition system environment to avoid reactions with water, oxygen or nitrogen compounds that may cause problems in later processing stages. Furthermore, annealing in the presence of hydrogen may help to increase the number of Si-Z-Si bonds that are formed between the silane end of the silane molecule and adjacent silicon-containing layers. Annealing also may help to improve the crystallinity of the low dielectric constant polymer layer. For example, as described above, the crystallinity of the polymer chains of the low dielectric constant polymer material may improved from 20-50% to about 70% during this annealing step.

Composite polymer dielectric film 10 may be annealed at any suitable temperature, and under any suitable concentration of hydrogen and/or mixture of hydrogen with other inert gases such as noble gases. Depending upon the temperature regime used, nitrogen may not be suitable for this process, as nitrogen may react with radicals in the composite film under annealing conditions. Examples of suitable annealing temperatures include, but are not limited to, temperatures sufficient to promote the reactions of free radicals in the composite polymer film with hydrogen, yet that are below the melting points of the low dielectric constant polymer layers. In the specific case of a composite film formed from an acrylic or methacrylic silane and PPX-F, annealing temperatures of between 300 and 400 degrees Celsius may be sufficient to eliminate free radicals within composite polymer dielectric film 10, and to increase the crystallinity of the PPX-F layer and to create a chemically, electrically and dimensionally stabilized film.

Composite polymer dielectric film 10 may be annealed under any suitable pressure and concentration of hydrogen. For example, composite polymer dielectric film 10 may be annealed under pure hydrogen gas (of any suitable purity), or may be annealed under hydrogen gas diluted in another inert gas. Examples of suitable diluent gases include, but are not limited to, inert gases such as argon or helium. The total pressure while annealing composite polymer dielectric film 10 may be between approximately 200 mTorr to 500 Torr, and more typically between approximately 1 to 20 Torr.

Likewise, composite polymer dielectric film 10 may be heated using any suitable heat source. For example, a hotplate may be used to heat composite polymer dielectric film 10 during annealing, or an indirect radiative heat source may be used.

Composite polymer electric film 114 may be annealed under hydrogen for any suitable amount of time. For example, in the specific case of a composite film formed from an acrylic or methacrylic silane and PPX-F annealed at a temperatures of between 300 and 400 degrees Celsius, an annealing time of between 2 and 10 minutes on a hot plate may be sufficient to eliminate free radicals and to increase the crystallinity of the PPX-F layer.

While the embodiment of Fig. 3 shows both the first silane layer and the second silane layer as being individually exposed to a free radical-generating energy source after the layers are deposited, it will be appreciated that polymerization of either the first silane layer, the second silane layer, or both the first and second silane layer may be initiated after all three layers have been deposited. In this instance, heat may be applied to the composite dielectric polymer film in the absence of hydrogen (and any free radical scavenging species) for a first interval to cause reaction with low dielectric constant layer 16 and, depending upon the chemical properties of the silane material, polymerization of these layers, and then heat may be applied for a second interval in the presence of hydrogen to eliminate any remaining free radicals.

As described above, even in the absence of first adhesion promoter layer 14, the adhesion of low dielectric constant polymer layer 16 may be improved by exposing the substrate to UV light for a brief interval before exposing the substrate to the diradical

monomer to create the low dielectric constant polymer layer. Fig. 4 shows, generally at 200, an embodiment of a method of forming a low dielectric constant polymer film in this manner. Method 200 includes first, at 202, exposing the substrate to UV light to remove water (and possibly other contaminants) from the surface of the substrate. The substrate
5 may be exposed to any suitable wavelength of UV light, and for any suitable interval. For example, the substrate may be exposed to UV light between 170 nm and 350 nm, or more specifically between 220 and 300 nm.

After exposing the substrate to UV light at 202, the substrate may next be exposed at 204 to a gas phase diradical monomer, such as one generated from the compounds
10 disclosed above in structures (V) and (VI), to form the low dielectric constant polymer layer. The substrate may be maintained in a high vacuum environment between exposure to the UV light at 202 and exposure to the gas phase diradical monomer at 204 to prevent the substrate surface from being re-contaminated by water or other contaminants. To maintain a sufficiently pure environment during deposition, the deposition system may be
15 configured to have a leakage rate of less than approximately 2 mTorr/minute, and more specifically less than about 0.4 mTorr/min. After completing the growth of the low dielectric constant polymer film at 204, the low dielectric constant polymer layer may be annealed in the presence of hydrogen, as described above in the context of Fig. 3.

Fig. 5 shows, generally at 300, a block diagram of an exemplary system for
20 forming composite dielectric polymer system 10. System 300 is described in the context of a system for depositing a PPX-based polymer, but it will be appreciated that system 300 may be adapted for use with any other suitable low dielectric constant polymer film.

System 300 includes one or more process modules 302 (a second process module is shown generally at 302a), a post-treatment module 304, a transfer module 306 connecting the process module 302 and the post-treatment module 304, an equipment front-end module 308 (EFEM) that interfaces system 300 with an exterior environment, and first and second load locks 310 and 312 for respectively introducing substrates into and removing substrates from transfer module 306.

System 300 also has a system controller 314 including memory 316 having instructions stored thereon that are executable by a processor 318 for controlling the various parts of system 300 to effect the formation of composite polymer dielectric film

10. Controller 314 is depicted in Fig. 5 as having electrical connections to a process module controller 314a on process module 302, a post-treatment controller 314b on post-treatment module 304, and to transfer module 306 (which may also include its own controller, not shown). However, it will be appreciated that controller 314 may have electrical connections to any other component having electrical systems, including but not limited to load lock 310, load lock 312, equipment front end module 308, and any sub-components contained within the general components shown in Fig. 5. The depicted system configuration allows all three layers of composite polymer dielectric film 10 to be deposited and annealed without breaking the deposition vacuum, and thus achieving high system throughputs while increasing manufacturing yields.

It will be appreciated that the electrical portion of Fig. 5 merely sets forth an exemplary hardware architecture, and does not indicate a particular physical relationship between the system controller and individual module controllers. Moreover, it will be

appreciated that the depicted hardware architecture is merely exemplary, and that system 10 may utilize any other suitable architecture. For example, a single controller may be used to control all functions of all modules, rather than having a controller for each module. Furthermore, while only system controller 314 is depicted as having memory and a processor, each of the individual module controllers typically also has memory and one or more processors associated therewith.

Process module 302 is configured to form first adhesion promoter layer 14, low dielectric constant polymer layer 16, and second adhesion promoter layer 18. Process module 302 includes a silane delivery system 319 with a silane container 320 and a carrier gas source 322 (typically nitrogen or other suitable inert gas) fluidically connected with the silane container via a mass flow controller 324. Process module 302 also includes a gas phase monomer delivery system 325, described in more detail below.

Silane delivery system 319 is configured to deliver a flow of a silane into a deposition chamber 326 for forming first adhesion promoter layer 14 and second adhesion promoter layer 18 on a substrate in the deposition chamber. In addition to mass flow controller 324, various valves (not shown) may be disposed between silane container 320 and deposition chamber 326 to regulate the flow of into the deposition chamber.

System controller 314 electronically communicates with process module controller 314a to start and finish the silane deposition process, while process module controller 314a directs process module 302 to perform the various functions and operations that produce layers 14, 16 and 18 of composite polymer dielectric film. For example, process

module controller 314a maintains the clean vacuum environment for deposition chamber 326 and reactor 338 via a vacuum pump system (not shown), controls the temperature distribution of the heating elements 332, delivers a desired amount of PPX precursor from precursor canister 334 into deposition chamber 326 through vapor flow controller 5 336 and reactor 338, and effects the UV lamp 330 to remove water from the substrate surface, etc.

To effect the silane deposition process, process module controller 314a maintains a desired gas pressure and mixture within the deposition chamber 326. For example, the base pressure within deposition chamber 326 may be held below 0.5 mTorr, and more specifically below 0.05 mTorr and during deposition of the silane at 200 mTorr to 500 10 Torr, preferably between 2 to 20 Torr. Alternatively, higher pressures may be employed.

Process module controller 314a may introduce silane precursor into deposition chamber 326 at any suitable rate. The rate and/or amount of silane that is introduced into 15 deposition chamber 326 may be controlled by controlling the temperature of the silane container (which affects the vapor pressure of gas phase silane in the container) and the flow rate of nitrogen into the deposition chamber 326. Examples of suitable silane temperatures and nitrogen flows include temperatures in the range of 10 and 50 degrees Celsius, and flow rates between 50 and 500 sccm, although silane temperatures and 20 nitrogen flow rates outside of these ranges may also be used. It will be appreciated that silane container 320 may include various heaters, temperature controllers, level detectors, and other components electrically coupled to and controlled by controller 314 that are not

shown in the high-level block diagram of Fig. 5. Likewise, other components shown in Fig. 5 may include various detectors, pumps, temperature control systems, etc. that have been omitted from Fig. 5 for purposes of clarity.

Deposition chamber 326 may include a temperature-controlled chuck 328 for
5 maintaining a substrate at a controlled temperature within the chamber. Temperature-controlled chuck 328 may be an electrostatic chuck, and may include a gas outlet configured to allow a desired pressure of a heat transfer gas to be maintained between the substrate and chuck surface, thus helping to maintain the substrate surface at a substantially uniform temperature. For example, maintaining helium at a pressure of up
10 to 10 psi between the substrate and chuck has been found to keep the temperature across the surface of the substrate within one degree Celsius of a desired target temperature. Furthermore, the use of an electrostatic chuck as temperature-controlled chuck 328 may provide sufficient holding force between the chuck and the substrate to keep the leak rate for helium below 0.4 sccm.

15 Deposition chamber 326 may also include a UV lamp 330 configured to illuminate a substrate positioned on temperature-controlled chuck 328. UV lamp 330 may be used both to remove water (and other contaminants) from a substrate surface prior to depositing any film layers on the substrate, and also may be used to cause the free radical polymerization of the adhesion promoter layers and the chemical bonding of the adhesion
20 promoter layers with the low dielectric polymer layer. Deposition chamber 326 may, alternatively or additionally, include heating elements 332 or a plasma source 333 for causing the polymerization of the silane layers. UV lamp 330 and/or heating elements

332 may be positioned outside of deposition chamber 326 and irradiate through a quartz (or other UV-transparent material) window provided in the deposition chamber. Alternatively, a suitable UV lamp may be positioned within the deposition chamber.

Deposition chamber 326 may be configured to have a low leakage rate and to hold
5 a high vacuum to help avoid the introduction of free radical scavenger species, such as oxygen and amino-containing compounds, during the various layer deposition and polymerization processes. For example, deposition chamber 326 may be configured to hold a vacuum of less than 0.01 mTorr during the deposition of the low dielectric constant polymer layer, and have a leakage rate of less than approximately 2
10 mTorr/minute, and in some embodiments, as low as 0.4 mTorr/minute or less.

Process module 302 also includes a gas phase monomer delivery system 325 for delivering a low dielectric constant polymer free radical intermediate to deposition chamber 326. Gas phase monomer delivery system 325 includes a precursor vessel 334 for holding a precursor material, a vapor flow controller 336 for regulating the flow of
15 vapor phase precursor out the vessel, and a reactor 338 for creating a gas phase free radical intermediate from the precursor. For example, where the precursor is the monomer $\text{CF}_2\text{BrC}_6\text{H}_4\text{CF}_2\text{Br}$, reactor 338 may convert the precursor to the diradical $^*\text{CF}_2\text{C}_6\text{H}_4\text{CF}_2^*$ for the formation of a PPX-F film. Furthermore, reactor 338 may be configured to chemically react with the bromine leaving groups and to be periodically
20 regenerated to remove bromine from the reactor. Examples of suitable reactors include those disclosed in U.S. Patent Application Serial No. 10/243,990 of Lee et al., filed September 13, 2002, the disclosure of which is incorporated by reference.

Process module controller 314a may be configured to maintain precursor vessel 334 at any suitable temperature for producing a desired vapor pressure of precursor monomer. For example, where $\text{BrCF}_2\text{C}_6\text{H}_4\text{CF}_2\text{Br}$ is the monomer precursor, process module controller 314a may maintain precursor vessel 334 at a temperature of between approximately 40 and 90 degrees Celsius to create a sufficient vapor pressure of the monomer. Likewise, process module controller 314a may be configured to maintain any suitable rate of flow of vapor phase precursor into reactor 338 via vapor phase controller 336. Examples of suitable flow rates of precursor into reactor 338 include, but are not limited to, flow rates of between 1 and 50 sccm, +/- 1 sccm.

As mentioned above, system 300 may include one or more process modules 302. While the embodiment depicted in Fig. 5 includes two process modules 302, it will be appreciated that system 300 could include three or more process modules if desired. The use of multiple process modules may help to increase substrate throughput. Alternatively, a designated process chamber for deposition of the first and second silane layers can also be employed where more than two process modules are included in a deposition system, as described in more detail below.

Post-treatment module 304 is configured to perform the above-described post-deposition heating and/or annealing processes. Furthermore, post-treatment module 304 may be configured to perform the deposition of first adhesion promoter layer 14, second adhesion promoter layer 18, or both adhesion promoter layers instead of (or in addition to) process modules 302 and 302a. Post-treatment module 304 includes an annealing gas source 340, at least one valve 342 to control a flow of annealing gas from the annealing

gas source 340 into a processing chamber 344, and one or more heating elements 346 for heating a substrate during an annealing process.

Processing chamber 344 may be configured to process substrates one at a time, or may be equipped for batch processing, with such features as a wafer elevator 348, etc., where equipped for batch processing, processing chamber 344 may include a plurality of heating elements 346 to provide consistent heating to each of a plurality of wafers being processed. The use of a wafer elevator 348 allows a plurality of substrates in processing chamber 344 to be vertically stacked, which may help to reduce the footprint of processing chamber 344 relative to a batch processor in which the substrates are not vertically stacked. The number of heating elements 346 and the height of wafer elevator 348 is determined at least in part by the number of process modules 302 used in system 300.

Processing chamber 344 of post-treatment module 304 may be configured to have a low leakage rate and be able to hold a high vacuum to help avoid the introduction of free radical scavengers such as O_2 and H_2O from outside of the chamber. Examples of suitable leak rates for processing chamber 344 of post-treatment module 304 include, but are not limited to, rates below about 2 mTorr/minute. Heating elements 346 of post-treatment module 304 may be radiant heaters, one or more hotplates on which substrates rest during processing, or may take any other suitable form. Heating elements 346 are controllable by controller 314 to heat a substrate to any suitable temperature for annealing. In the specific case of a PPX-F low dielectric constant polymer film, the

heating elements may be configured to heat the substrates to a constant controlled temperature of between approximately 300-450 (+/- 2-5) degrees Celsius.

Post-treatment module 304 may also be configured to deposit first adhesion promoter layer 14 and/or second adhesion promoter layer 318. In this instance, post-treatment module 304 may include a silane precursor delivery system 350 configured to deliver a flow of silane precursor into processing chamber 344 for forming first adhesion promoter layer 14 and second adhesion promoter layer 18. Silane precursor delivery system 350 may be similar in structure and function to silane precursor delivery system 319 described above in the context of process module 302. Post-treatment module 304 also may include a UV lamp (or other free radical-generating energy source, such as a plasma source) for polymerizing the silane precursor.

Post-treatment module controller 314b directs post-treatment module 304 to perform the various functions and operations that go into the post-deposition processing of composite polymer dielectric film 10. For example, post-treatment module controller 314b regulates the composite film annealing temperature through heating elements 346, maintains prescribed hydrogen/helium pressure within processing chamber 344 via the gas source 340 and controlling valve 342, and controls the vertical indexing motion of the wafer elevator 348 for wafer transfer into and out of post-treatment module 304. System controller 314 may communicate with post-treatment module controller 314b to cause the post-treatment module controller to start and stop these processes.

Transfer module 306, equipment front end module 308, load lock 310 and load lock 312 each may be considered part of a platform that supports use of process

module(s) 302 and post-treatment module 304. During a deposition process, a substrate is loaded initially into equipment front-end module 308, where alignment systems within the equipment front-end module pre-align the substrate. The substrate is then moved via a mechanical arm or other transfer apparatus (not shown in Fig. 5) within equipment front-end module 308 to load lock 310, which is pumped down to a pressure of a few mTorr. From load lock 310, another substrate transfer apparatus (not shown in Fig. 5) within transfer module 306 transfers the substrate from load lock 310 into the transfer module, and then into the silane deposition module for an initial exposure to UV light and/or deposition of the first adhesion promoter layer.

Transfer module 306 is also controllable by controller 314 to transfer a substrate among process module(s) 302 and post-treatment module 304 at appropriate times during the manufacture of composite dielectric polymer film 10. To help prevent reaction of unreacted free radicals in composite dielectric film with free radical scavengers while the substrate is being transferred between process module(s) 302 and post-treatment module 304, transfer module 306 may be maintained at a suitable level of vacuum. Suitable levels of vacuum include, but are not limited to, pressures below 0.1 mTorr. Maintaining transfer module 306 at pressures in this range may help to avoid the introduction of harmful quantities of free radical scavengers into transfer module 306, and thus may help to prevent unwanted reactions.

Once all processing steps have been completed, transfer module 306 may be controlled by controller 314 to transfer a substrate into load lock 312, where it may be

brought back to the pressure of equipment front-end module 308 for removal from system 300.

Fig. 6 shows, generally at 400, a block diagram of another exemplary system for forming composite dielectric polymer system 10. Like system 300, system 400 includes one or more process modules 402 (a second process module is shown generally at 402a), a post-treatment module 404, a transfer module 406 connecting the process module 402 and the post-treatment module 404, an equipment front-end module 408 (EFEM) that interfaces system 400 with an exterior environment, and first and second load locks 410 and 412 for respectively introducing substrates into and removing substrates from transfer module 406. System 400 also includes a system controller 414 including memory 416 having instructions stored thereon that are executable by a processor 418 to control the various functions of system 400. System controller 414 may interface with separate module controllers 414a, 414b, which control the individual functions of each module.

System 400 also includes a separate silane deposition module 420 for depositing layers 14 and 18 of composite polymer dielectric film 10. Silane deposition module 420 includes a silane deposition chamber 422 to which a silane delivery system 424 is connected. Silane delivery system 424 is configured to deliver a flow of a silane into silane deposition chamber 422 for forming first adhesion promoter layer 14 and second adhesion promoter layer 18 on a substrate positioned in the silane deposition chamber. Silane delivery system 424 includes a silane container 426 and a carrier gas source 428 (typically nitrogen or other suitable inert gas) fluidically connected with the silane

container via a mass flow controller 430. Other valves may be disposed between silane container 426 and silane deposition chamber 422 preventing or allowing silane to flow into the silane deposition chamber.

Silane deposition chamber 422 may include a free radical-generating energy source for creating free radicals in the silane molecules to allow the polymerization of the silane molecules, and reactions between the silane molecules and the low dielectric constant polymer layer. One example of a suitable free radical generating source is a UV lamp 425. UV lamp 425 may also be used to remove water from substrate surfaces prior to depositing layer 14 of composite polymer dielectric film 10. Alternatively, another energy source, such as a plasma source 427 or a thermal energy source, may be used.

Silane deposition module 420 also includes a silane deposition module controller 414c for controlling the various processes and operations of the silane deposition process, including but not limited to the delivery of silane precursor from the silane canister to the deposition chamber, and the temperature of temperature controlled chuck. System controller 414 electronically communicates with silane deposition module controller 414c to start and finish the silane deposition process. To effect the silane deposition process, silane deposition module controller maintains the gas pressure of the silane deposition chamber, the temperature of the silane precursor, the flow of the carrier gas, etc. For example, the base pressure within silane deposition chamber may be held below 0.5 mTorr, and more specifically below 0.05 mTorr. During deposition of the silane, pressure within silane deposition chamber 424 may be held at a pressure of between

approximately 200 mTorr and 500 Torrs, and more specifically between 2 to 20 Torrs. Alternatively, higher pressures may be employed.

Silane deposition module controller 414c may be configured to introduce silane into silane deposition chamber at any suitable rate. Examples of suitable silane temperatures and nitrogen flows include temperatures in the range of 10 and 50 degrees Celsius, and flow rates between 50 and 500 sccm, although silane temperatures and nitrogen flow rates outside of these ranges may also be used. It will be appreciated that silane container 320 may include various heaters, temperature controllers, level detectors, and other components electrically coupled to and controlled by silane deposition module controller that are not shown in the high-level block diagram of Fig. 6. Likewise, other components shown in Fig. 6 may include various detectors, pumps, temperature control systems, etc. that have been omitted from Fig. 6 for purposes of clarity.

While the depicted silane deposition module includes only a single silane delivery system, it will be appreciated that the silane deposition module may include plural silane delivery systems. For example, where first adhesion promoter layer 14 is made of a silane having the general formula $(\text{ETO})_3\text{-Si-(WT)}$ while third adhesion promoter layer 18 is made of a silane having the general formula $(\text{ETO})\text{-Si-(WT)}_3$, the silane deposition module may have a separate silane delivery system for each of these silanes.

Fig. 7 shows, generally at 500, a plan view of an embodiment of a processing system for forming composite dielectric film 10 arranged on a fabrication room floor. System 500 includes three process modules 502, one post-treatment module 504, and one silane deposition module 520. A transfer module 506 is disposed in a central location,

and the process modules 502, post-treatment module 504, and load locks 310 and 312 are disposed around the transfer module. Transfer module 506 includes a pivotal mechanical arm 508 configured to transfer a wafer between wafer conveyors 522 and 524 on load locks 510 and 512, respectively, wafer conveyors 526, 528 and 530 on process modules 502, and wafer conveyor 532 on post-treatment module 304. The depicted configuration of system 500 allows a substrate to be transferred between load locks 510 and 512, transfer module 506, process modules 502, post-treatment module 504 without breaking the overall system vacuum. Substrates may be sequentially processed in process modules 502, and then annealed in post-treatment module 504 in a batch manner for a high system throughput. Using the above-disclosed processing equipment and conditions, composite polymer dielectric film 10 can be deposited and annealed on a wafer in 4 to 6 minutes, which is within the time tolerances dictated by the overall fabrication speed of typical integrated circuit processing lines. It will be appreciated that the configuration shown in Fig. 7 is merely exemplary, and that any other suitable arrangement of the various components of systems 300 and/or 400 may be used.

Although the present disclosure includes specific embodiments of various composite dielectric films, methods of forming the films, and systems for forming the films, specific embodiments are not to be considered in a limiting sense, because numerous variations are possible. The subject matter of the present disclosure includes all novel and nonobvious combinations and subcombinations of the various films, processing systems, processing methods and other elements, features, functions, and/or properties disclosed herein. The following claims particularly point out certain

combinations and subcombinations regarded as novel and nonobvious. These claims may refer to “an” element or “a first” element or the equivalent thereof. Such claims should be understood to include incorporation of one or more such elements, neither requiring nor excluding two or more such elements. Other combinations and subcombinations of features, functions, elements, and/or properties may be claimed through amendment of the present claims or through presentation of new claims in this or a related application. Such claims, whether broader, narrower, equal, or different in scope to the original claims, also are regarded as included within the subject matter of the present disclosure.